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Description

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## Mixtures of finely ground waxes

The invention relates to mixtures of finely ground waxes, comprising two or more components A, B, C, and/or D, and to their use.

The use of waxes as processing aids for plastics, for the dispersing of pigments in plastics, as additive in printing inks and in coatings, as a processing aid for powder coatings, and in a number of other applications, is known. Many of these applications need waxes in ground form, permitting lower energy usage in processing, and therefore more cost-effective processing, and better dispersion, and smaller amounts added. Products of this type based on hydrocarbon waxes or on amide waxes are known. These are easy to grind and, for various applications, are also combined with other products, e.g. PTFE powders.

The compatibility of these non-polar waxes with non-polar media, such as aliphatic or aromatic solvents, polyethylene, polypropylene, and other non-polar substances, is very good. However, there is an increasing need for finely ground waxes for polar systems, too, because in this case non-polar waxes cannot always be used, because the result is often incompatibility.

Alternatives here are oxidized hydrocarbon waxes or polar natural waxes and their derivatives, such as montane wax acid derivatives.

Products of this type are known, examples being <sup>®</sup>Ceridust 121 or <sup>®</sup>Ceridust E/OP from Clariant GmbH, but their application is subject to restrictions, because grinding yield is low and this is associated with poor cost-effectiveness. As an alternative, wax powders can be produced by spray drying, but here there are restrictions in terms of heat resistance and viscosity, and also in the combination of various waxes or wax powders.

An object was therefore to find a technically simple method of providing finely ground polar waxes, so that the performance advantages of fine waxes can also be utilized in applications in polar media.

This object is achieved via mixtures of finely ground waxes, comprising two

or more components A, B, C, and/or D, wherein ester waxes are used as component A, amide waxes are used as component B, hydrocarbon waxes are used as component C, and oxidized long-chain hydrocarbon waxes are used as component D.

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The ester waxes are preferably natural ester waxes are synthetic ester waxes.

The natural ester waxes are preferably montane waxes, carnauba wax, 10 candelilla wax, and/or sugarcane wax.

It is preferable that the montane waxes are montane wax acid, derivatives of montane wax acid, e.g. esters of montane wax acid, soaps of montane wax acid, esteramides of montane wax acid, and/or a mixture of derivatives of montane wax acid with long-chain fatty acids.

The synthetic ester waxes are preferably sorbitan esters of unsaturated fatty acids, esters of polyols, such as pentaerythritol, glycerol, trimethylolpropane, with long-chain fatty acids, and/or their mixtures.

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Other preferred synthetic ester waxes are copolymers composed of longchain olefins having from 5 to 18 carbon atoms and of unsaturated acids, such as acrylic acid, methacrylic acid, maleic anhydride, itaconic acid, and/or derivatives of these acids.

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The derivatives are preferably methyl, ethyl, butyl, and/or 2-ethyhexyl esters of acrylic acid, methacrylic acid, maleic anhydride, and/or itaconic acid.

The copolymers are preferably those whose carboxy functionality has been modified via reaction with long-chain alcohols, with perfluoroalkyl alcohols, with short-chain amines, and/or with long-chain amines.

The long-chain alcohols are preferably tallow fatty alcohol, coconut fatty alcohol, and/or oxo alcohols.

The perfluoroalkyl alcohols are preferably C<sub>8-</sub>C<sub>18</sub>-perfluoroalkylpropanol, and/or narrower cuts of these alcohols.

Examples of the classes of these perfluoroalkylpropanols are  $C_8$ - $C_{12}$ -,  $C_{14}$ - $C_{18}$ -,  $C_8$ - $C_{14}$ -, and other perfluoroalkylpropanols with a different number and distribution of carbon atoms.

5 The short-chain amines are preferably butylamine, dimethylaminopropylamine, diethylaminoethanol, tetramethylpiperidinol, and/or diacetonediamine.

The long-chain amines are preferably tallow fatty amine, octylamine, 10 palmitylamine, and/or stearylamine.

The amide waxes are preferably reaction products composed of an amine component and of long-chain fatty acids or hydroxy fatty acids, and/or their mixtures.

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The amine components are preferably ethylenediamine and/or hexamethylenediamine.

The amide waxes are preferably reaction products composed of ammonia and of long-chain fatty acids or hydroxy fatty acids, and/or their mixtures.

The amide waxes are preferably reaction products composed of long-chain amines, such as hydrogenated tallow fatty amine, stearylamine, palmitylamine, coconut fatty amine, and of long-chain fatty acids or hydroxy fatty acids, and/or their mixtures.

The hydrocarbon waxes are preferably polyethylene waxes.

Preference is given to those polyethylene waxes which have been prepared by the Ziegler process or by means of metallocene technology.

As an alternative, the polyethylene waxes are Fischer-Tropsch waxes.

The inventive mixture preferably comprises the following finely ground 35 waxes:

from 40 to 99% by weight of component A from 0 to 95% by weight of component B from 0 to 95% by weight of component C from 0 to 95% by weight of component D,

where the entirety of the components is 100% by weight.

The inventive mixture particularly preferably comprises the following finely ground waxes:

from 40 to 90% by weight of component A
from 0 to 60% by weight of component B
from 0 to 60% by weight of component C
from 0 to 60% by weight of component D,
where the entirety of the components is 100% by weight.

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The invention also provides the use of the inventive mixtures of finely ground waxes as additive in printing inks, or in powder coatings or other coatings.

The invention also provides the use of inventive mixtures of finely ground waxes for the dispersion of pigments and of additives in plastics.

The invention also provides the use of inventive mixtures of finely ground waxes as hydrophobicizing additive in crop-protection preparations.

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Finally, the invention also provides the use of mixtures of finely ground waxes as lubricant in various plastics.

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Approaches to the improvement of grinding technology for the pure components did not lead to the desired result. Despite optimized technology, the grinding yields were higher with the inventive wax combinations than with the pure products.

Component	Compound	Example 1 (comparison)	Example 2 (inventive)	Example 3 (inventive)
Α	Glycerol monotanate	85	85	85
Α	Montane wax acid	15	15	15
В	Amide wax C	-	10	
С	PE wax PE 130		-	10
	Mixture	100	110	110
	Grinding performance g/h	350	800	500

## Example mixture 2

Component	Compound	Example 4 (comparison)	Example 5	Example 6 (inventive)
			(inventive)	
Α	Pentaerythritol monotanate	85	85	85
Α	Montane wax acid	15	15	15
В	Amide wax C	-	15	-
С	PE wax PE 130	-	-	15
	Mixture	100	115	115
	Grinding performance g/h	350	500	450

## Example mixture 3

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Component	Compound	Example 8	Example 9	Example 10
A	Ethanediol	10	10	10
·	monomontanate			
Α	Ethanediol	20	20	20
	dimontanate			
Α	Butanediol	20	20	20
	1,3-montanate			
	Ca montanate	45	45	45
Α	Montane wax acid	5	5	5
В	Amide wax C	-	10	_
С	PE wax PE 130	_		15
	Mixture	100	115	115
	Grinding	350	500	450
	performance g/h			

Component	Compound	Example 11	Example 12	Example 13
A	Sorbitol monotanate	85	85	85
Α	Montane wax acid	15	15	15
В	Amide wax C	-	20	
С	PE wax PE 130	-	_	35
	Mixture	100	120	135
	Grinding	450	650	600
	performance g/h			

Component	Compound	Example 14	Example 15
·		(comparison)	(inventive)
Α	Sorbitol montanate/stearate	100	85
В	Amide wax HS	-	15
	Mixture	100	100
	Grinding performance g/h	500	800

Component	Compound	Example 16	Example 17
<u>'</u>		(comparison)	(inventive)
Α	Carnauba wax	100	85
С	PE wax H2	-	15
	Mixture	100	100
	Grinding performance g/h	350	550

Component	Compound	Example 18 (comparison)	Example 19 (inventive)
Α	Modified olefin copolymer	100	85
В	Amide wax	<u>-</u>	15
	Mixture	100	100
	Grinding performance g/h	1000	1600

Component	Compound	Example 20	Example 21
·	·	(comparison)	(inventive)
Α	Olefin copolymer CE 1	100	85
С	PE wax	-	15
	Mixture	100	100
	Grinding performance g/h	400	650

Component	Compound	Example 22	Example 23
•		(comparison)	(inventive)
Α	Montane wax acid	0	5
D	PE wax oxidate	100	95
	Mixture	100	100
	Grinding performance g/h	400	650

The grinding process used an AFG 100 countercurrent fluidized-bed mill, Hosokawa Alpine. The target size for the particles was D $_{50}$  of 10-12  $\mu m$ .

The examples lists show that the inventive mixtures of finely ground waxes give a considerable improvement during the grinding process.

## Application examples:

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Pigment dispersion in powder coating:

It is known that the dispersion of the pigments can be improved via the use of montane waxes. In order to achieve a high level of activity it is necessary that the products are used in the form of powders or micropowders. It has now been found that use of the inventive mixtures can achieve not only improved pigment dispersion but also an increase in extrusion output. At the same time, the costs for the production of the inventive wax combination were markedly lower when comparison is made with a pure montane wax, and therefore operations with the inventive wax mixtures can be substantially more cost-effective.

Examples of formulations for powder coating (all data in parts by weight) Power coating A:

70.0 parts of

binder

20 9.0 parts of

Blanc Fixe N

20.0 parts of

®Kronos 2310 titanium dioxide

1.0 part of

®Hostaperm Blue A4R

0.5/1.0/2.0 part(s) of

montanic acid derivative (<sup>®</sup>Ceridust 5551, Clariant

GmbH)

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Power coating B:

70.0 parts of

binder

9.0 parts of

Blanc Fixe N

20.0 parts of

<sup>®</sup>Kronos 2310 titanium dioxide

30 1.0 part of

®Red Violet ER 02

0.5/1.0/2.0 part(s) of

montanic acid derivative (<sup>®</sup>Ceridust 5551, Clariant

GmbH)

Power coating C:

35 70.0 parts of

binder

9.0 parts of

Blanc Fixe N

20.0 parts of

®Kronos 2310 titanium dioxide

1.0 part of

®Hostaperm Blue A4R

1.0 part of

waxes of example 4/example 2/example 3

Power coating D:

70.0 parts of

binder

9.0 parts of

Blanc Fixe N

5 20.0 parts of

®Kronos 2310 titanium dioxide

1.0 part of

®Red Violet ER 02

1.0 part of

waxes of example 4/example 2/example 3

were produced by way of the following steps in a process

10 - premixing all the components in a Mixaco mixer

extrusion in a twin-screw APV Baker laboratory extruder at 110°C

- comminuting in a Retsch pinned disc mill

- sieving to a grain fineness smaller than 125 μm

15 The composition of the binder is:

90.13%

<sup>®</sup>Alftalat AN 989 (Vianova Resins)

4.74%

<sup>®</sup>Primid XL 552 (EMS Chemie)

4.37%

<sup>®</sup>Additol XL 9824 (Vianova Resins)

0.29%

® Benzoin

20 0.47%

®Hostanox M 101 (Clariant GmbH)

The average particle size after extrusion was 9  $\mu\text{m}.$ 

The powder was applied by means of an electrostatic spraying device from Wagner to metal substrates and stoved for 10 minutes at 180°C. The color strength of the coating is measured to DIN 55986 in a CM 3600d spectrophotometer from Minolta. To calculate the relative color strength, the measured value is based on the value measured for the control specimen (without dispersing agent; control specimen = 100%). Values greater than 100 mean higher color strength than the control specimen, and values smaller than 100 mean lower color strength.

Inventive wax	Hostaperm Red Violet	Hostaperm Blue A4R
	ER 02	
Parts by weight	Rel. color strength %	Rel. color strength %
0	100	100
0.5	.108	114
1.0	120	123
2.0	127	132

Inventive wax	Hostaperm Red Violet ER 02	Hostaperm Blue A4R
	EN UZ	
1.0 part by weight	Rel. color strength %	Rel. color strength %
None	100	100
Ceridust 5551	120	123
Example 4	119	125
Example 2	123	120
Example 3	121	122

It could be shown that the addition of an inventive mixture as dispersing agent improves the dispersion of pigments and therefore increases the color strength.